

Application No.: 09/815,311  
Attorney Docket No.: 05788.0157-00

**CLAIM OF FOREIGN PRIORITY**

Applicants have claimed priority based on EP98118194.4. A certified copy of the priority application was filed on March 23, 2001. To date, the Examiner has not acknowledged the claim for priority and submission of this priority document. The Examiner is respectfully requested to acknowledge the claim of priority and receipt of the certified copy in the next Communication from the Office.

**REJECTION UNDER 35 U.S.C. § 102(b)**

In the Final Office Action dated November 14, 2002, the Office has maintained the rejection of claims 16-30 under 35 U.S.C. § 102(b)<sup>1</sup> as anticipated by *Sonoda et al.* (U.S. Patent No. 5,707,732) in view of *Betso et al.* (U.S. Patent No. 6,262,161) to show the state of the art. Final Office Action at 2-3. Applicants traverse this rejection for at least the reasons of record and the following additional reasons.

Applicants submit that *Sonoda et al.* does not disclose each of the limitations of Applicants' claimed invention. M.P.E.P. § 2131; see also *Electro Med. Sys., S.A., v. Cooper Life Sciences, Inc.*, 32 U.S.P.Q.2d 1017, 1019-20 (Fed. Cir. 1994). For example, *Sonoda et al.* fails to disclose, expressly or inherently, (1) "natural magnesium" and (2) "hydrolyzable organic silane groups . . . for compatibilization of the natural magnesium hydroxide" with the polymeric components or polymeric matrix, as recited by Applicants' claims.

<sup>1</sup> While the rejection has been stylized as a rejection under §102(b), in view of Applicants' claim of benefit to Provisional Application No. 60/102,926, Applicants believe the rejection should have been made under §102(e). Regardless, the pending claims are not anticipated by *Sonoda et al.* under either section of 102 for substantially the same reasons.

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First, it appears as though the Office misunderstands Applicants' invention, particularly with respect to the disclosure of *Sonoda et al.* Applicants' invention relates, *inter alia*, to the use of natural magnesium hydroxide. As noted on page 6 of Applicants' specification, Applicants discovered that "[t]here are remarkable advantages when using natural magnesium hydroxide." In contrast, synthetic magnesium hydroxide "has a considerable impact on the cost of the finished product, so as to make flame-retardant systems based on magnesium hydroxide non-competitive when compared with the halogen-containing flame-retardant compositions described above." Specification at 3, lines 18-22. Natural magnesium hydroxide is "obtained by grinding minerals based on magnesium hydroxide, such as brucite." *Id.* at 12, lines 1-2. Accordingly, natural magnesium hydroxide "generally contains various impurities." *Id.* at 12, lines 22. Furthermore, "natural magnesium hydroxide has a highly irregular granular morphology in terms both of its geometric shape and of its surface appearance. In contrast, the magnesium hydroxide obtained by precipitation consists of flattened hexagonal crystallites that are substantially uniform both in size and morphology." *Id.* at 13, lines 20-26. Thus, natural magnesium hydroxide is not identical to synthetic magnesium hydroxide.

*Sonoda et al.*, despite the Office's assertions to the contrary (Final Office Action at 3), does not disclose natural magnesium hydroxide obtained by grinding minerals.

Instead, it states:

As hydrated inorganic flame retardant fillers, magnesium hydroxide (preferred) or alumina trihydrate are used. While conventional off-the-shelf magnesium hydroxide and alumina trihydrate can be used, a preferred magnesium hydroxide has the following characteristics: (a) a strain in the <101> direction of no more than  $3.0 \times 10^{-3}$ ; (b) a crystallite size in the <101> direction of more than

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800 angstroms; and (c) a surface area, determined by the BET method, of less than 20 square meters per gram. The preferred magnesium hydroxide and a method for its preparation are disclosed in U.S. Pat. No. 4,098,762. A preferred characteristic of this magnesium hydroxide is that the surface area, as determined by the BET method, is less than 10 square meters per gram.

Sonoda et al., col. 6, line 60-col. 7, line 5. Notably, the preferred magnesium hydroxide of Sonoda et al., which is the only magnesium hydroxide disclosed with any detail whatsoever, specifies a crystallite size. As noted by the distinction between natural and synthetic magnesium hydroxide in Applicants' disclosure at page 13 (and as mentioned above), this means that Sonoda et al. discloses synthetic magnesium hydroxide. This is further highlighted by Sonoda et al.'s stated preference for magnesium hydroxide as disclosed in U.S. Patent No. 4,098,762 (courtesy copy attached), which concerns a synthetic magnesium hydroxide. Thus, Sonoda et al. fails to disclose the limitation of "natural magnesium hydroxide."

Furthermore, it would not have been obvious at the time the invention was made to modify the teachings of Sonoda et al. with a reasonable expectation of success in creating every element of Applicants' claimed invention. For example, substituting natural magnesium hydroxide for synthetic magnesium hydroxide would not be obvious in light of the "research efforts . . . directed towards modifying properties of magnesium hydroxide to improve its compatibility with the polymer matrix and its degree of purity." Specification at 3, lines 1-4.

Second, the Office incorrectly assumes that the disclosure by Sonoda et al. of the grafting a copolymer with an alkenyl trialkoxy silane in the presence of an organic peroxide (col. 6, lines 41-46) is an inherent disclosure of "hydrolyzable organic silane groups . . . for compatibilization of the natural magnesium hydroxide" with the polymeric

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components or polymeric matrix, as recited by Applicants' claims. Final Office Action at 3-4.

Inherency, however, requires that "the missing descriptive matter is necessarily present in the thing described in the reference, and that it would be so recognized by persons of ordinary skill. Inherency . . . may not be established by probabilities or possibilities. The mere fact that a certain thing may result from a given set of circumstances is not sufficient." *In re Robertson*, 49 U.S.P.Q.2d 1949, 1950-51 (Fed. Cir. 1999) (quotation omitted) (emphasis added). The claim limitation must necessarily flow from the teaching of the prior art. *Ex parte Levy*, 17 U.S.P.Q.2d 1461, 1464 (Bd. Pat. App. & Int. 1990). Thus, for Sonoda et al. to anticipate Applicants' claims, Sonoda et al. must indisputably, not merely likely, disclose "hydrolyzable organic silane groups . . . for the compatibilization of natural magnesium hydroxide." It does not.

While Sonoda et al. admittedly discloses the use of a silane compound, that disclosure is merely an afterthought. See col. 6, lines 40-55. Sonoda et al. provides no examples of compositions utilizing such a compound and offers merely a cursory direction to one skilled in the art as to its use. Accordingly, the value of Sonoda et al. as an anticipating reference is highly questionable.

Furthermore, there is no basis for one skilled in the art to conclude that any of Sonoda et al.'s hydrolyzable organic silane groups compatibilize natural magnesium hydroxide, let alone synthetic magnesium hydroxide. Sonoda et al. requires that at least one of its polymers (i) and (ii) be modified with an anhydride of an unsaturated aliphatic diacid. Col. 2, lines 1-3. As discussed in depth in Applicants' specification, the sole purpose of Sonoda et al.'s anhydride is to compatibilize the filler. See, e.g.,

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specification at 6, lines 7-14. This limited use is further supported by the teachings of the prior art, which states that "maleic anhydride resins can be used as coupling agents between polymers -- mainly polyolefins such as polyethylene and polypropylene -- and fillers, in order to increase the filler acceptability of polymers." *Dupont Industrial Polymers: Fusabond®*, <http://www.dupont.com/industrial-polymers/fusabond/H-81624/H-81624.html> (courtesy copy attached). While Applicants cannot be certain that Fusabond® is the very same product used by Sonoda et al., the reference shows that one purpose of a grafted anhydride is provide resins with an affinity with flame-retardant fillers, such as magnesium hydroxide, in order to increase the filler acceptability of polymers.

Since Sonoda et al.'s anhydride grafted to the polymers already performs the very function Applicants' hydrolyzable organic silane groups perform, Sonoda et al.'s silane compounds are NOT present for compatibilization but rather are present for a wholly separate and different purpose, i.e., moisture curing of the polymers. Col. 6, line 42. Further, while Belso may disclose the silane compounds can act as couplers, that disclosure is negated by Sonoda et al.'s use of anhydrides to compatibilize the filler. Sonoda et al., therefore, does not inherently disclose "hydrolyzable organic silane groups . . . for the compatibilization of natural magnesium hydroxide."

Because Sonoda et al. fails to teach every limitation recited in those claims, independent claims 16, 29, and 30 are in condition for allowance. Claims 17-28 are in condition for allowance, for at least the reason that they depend from allowable claim 16, and thus are not anticipated by Sonoda et al.

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**CONCLUSION**

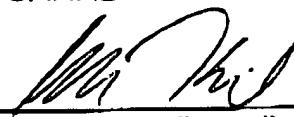
In view of the foregoing remarks, Applicants respectfully submit that this claimed invention is neither anticipated nor rendered obvious in view of the prior art references cited by the Office. Applicants, therefore, request the Examiner's reconsideration and reexamination of the application, and the timely allowance of the pending claims.

Please grant any extensions of time required to enter this response and charge any additional required fees to Deposit Account No. 06-0916.

Respectfully submitted,

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Dated: January 24, 2003

By: 

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Enclosures:

- (1) U.S. Patent No. 4,098,762
- (2) *Dupont Industrial Polymers: Fusabond®*,  
<http://www.dupont.com/industrial-polymers/fusabond/H-81624/H-81624.html>.

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## United States Patent [19]

Miyata et al.

[11] 4,098,762  
 [45] Jul 4, 1978

[54] MAGNETUM HYDROXIDES HAVING NOVEL STRUCTURE, PROCESS FOR PRODUCTION THEREOF, AND RESIN COMPOSITIONS CONTAINING THEM

[75] Inventors: Shigeo Miyata, Takamatsu; Masataka Kuroda, Kagawa; Akira Okada; Tomiharu Ohazaki, both of Takamatsu; Mitroo Takemoto, Marugame, all of Japan

[73] Assignee: Kyowa Chemical Industry Co., Ltd., Tokyo, Japan

[21] Appl. No.: 690,770

[22] Filed: May 27, 1976

[30] Foreign Application Priority Data

May 30, 1975 [JP] Japan 50-65049  
 Mar. 25, 1976 [JP] Japan 51-31919

[51] Int. Cl. 2 C08K 3/22; C08K 9/04  
 [52] U.S. Cl. 260/45.7 R; 106/306

[58] Field of Search 106/306, 308 S, 308 F;  
 260/45.7 R

[56]

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3,941,610	3/1976	Meekal .....	106/306 X

Primary Examiner—Sandra M. Person  
 Attorney, Agent, or Firm—Wenderoth, Lind & Ponack

[57]

## ABSTRACT

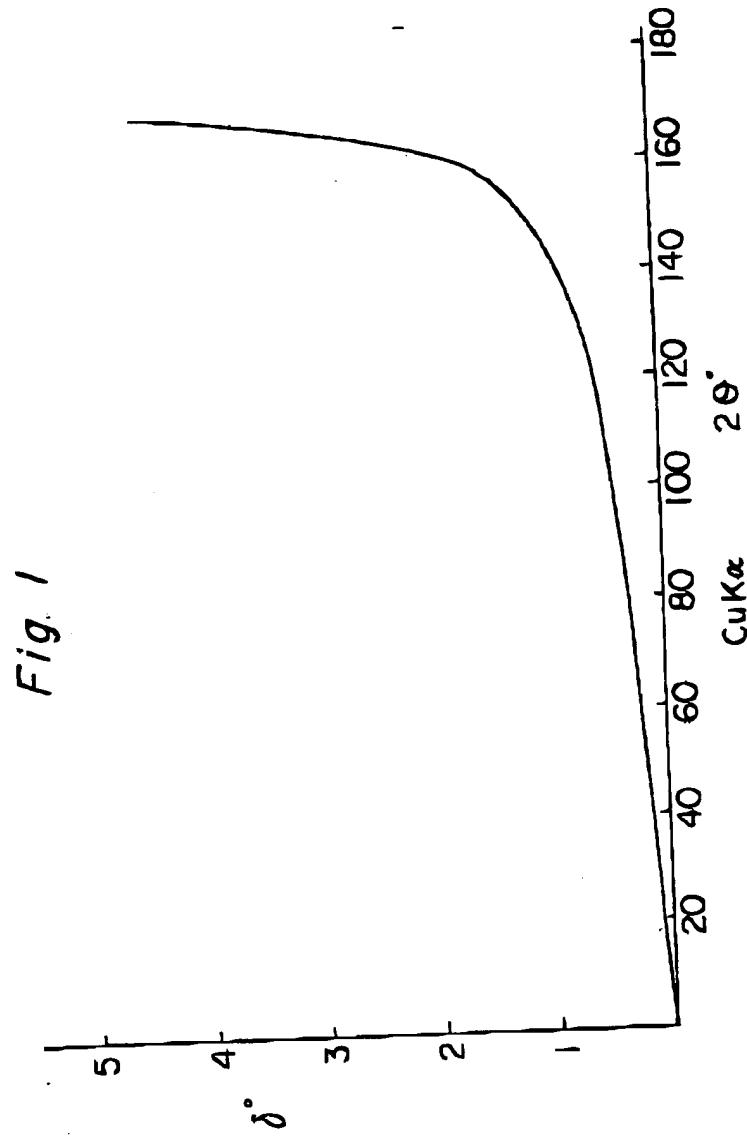
A novel magnesium hydroxide of the formula  $Mg(OH)_2 \cdot A_x \cdot mH_2O$  wherein A is Cl or  $NO_3$ , x is a number of more than 0 but less than 0.2, and m is a number of 0 to 6. The magnesium hydroxide, if desired as coated with anionic surfactants, is useful, for example, as fire retardants for thermoplastic synthetic resins and aqueous paints. The novel magnesium hydroxide is prepared by heating in an aqueous medium at an elevated pressure a novel basic magnesium chloride or nitrate of this invention expressed by the formula  $Mg(OH)_2 \cdot A_x \cdot mH_2O$  wherein A is Cl or  $NO_3$ , x is a number of more than 0 but less than 0.2, and m is a number of 0 to 6.

3 Claims, 3 Drawing Figures

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Fig. 3

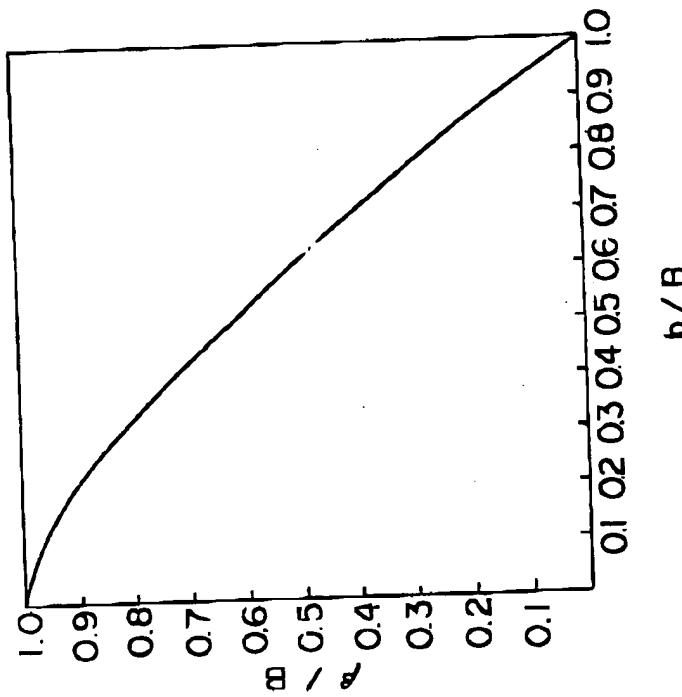
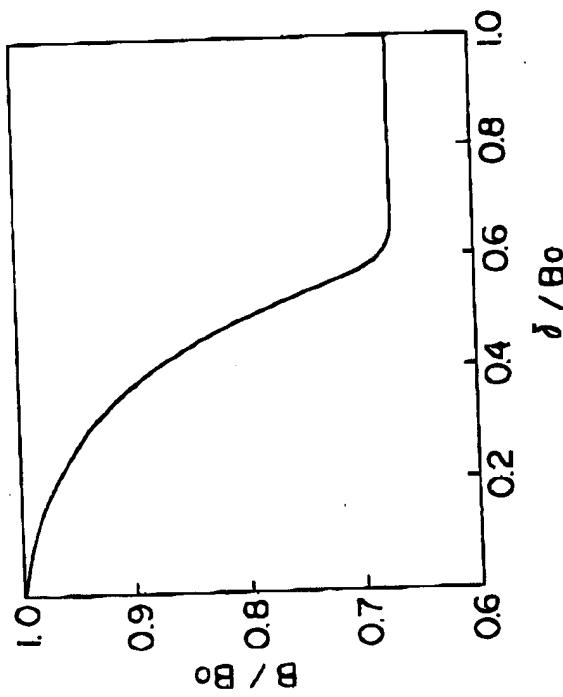


Fig. 2



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MAGNESIUM HYDROXIDES HAVING NOVEL  
STRUCTURE, PROCESS FOR PRODUCTION  
THEREOF, AND RESIN COMPOSITIONS  
CONTAINING THEM

This invention relates to magnesium hydroxides, coated or uncoated with anionic surface active agents, which have a novel X-ray diffraction pattern different from those of known magnesium hydroxides  $Mg(OH)_2$ , and exhibit superior operations and effects in such uses as fire retardants for thermoplastic synthetic resins and aqueous paints, or precursors of magnesium oxide for an annealing-separator for ferrosilicon, as compared with conventional available magnesium hydroxides; novel basic magnesium chloride or basic magnesium nitrate as intermediates thereof; processes for producing them, and to their use.

More specifically, the invention relates to a magnesium hydroxide, uncoated or coated with an anionic surface active agent, expressed by the following formula



which has a strain in the  $<101>$  direction of not more than  $3.0 \times 10^{-3}$ , a crystallite size in the same direction of more than 800 Å, and a specific surface area, determined by the BET method, of less than  $20 \text{ m}^2/\text{g}$ ; basic magnesium chloride or magnesium nitrate as its intermediate; processes for their preparation; and to their use, especially to a thermoplastic synthetic resin composition having improved properties such as superior fire retardancy or melt-shapability.

Magnesium hydroxide has been known from old, and used in a wide range of fields. For example, it is used to impart fire retardancy to thermoplastic synthetic resins. When it is incorporated in thermoplastic synthetic resins in an amount sufficient to impart a feasible fire-retarding effect to the resins, the physical properties of the resin, especially, impact strength or elongation, are deteriorated. Furthermore, the flow of the resin is reduced at the time of melt-shaping the resulting resin composition, and its shapability and shaping efficiency are reduced. Also, the resulting shaped articles frequently have a silver pattern which causes a poor appearance.

Our investigations led to the discovery that these defects are attributed to the structural characteristics which the conventional available magnesium hydroxides inherently have, especially to their strain, crystallite size and specific surface area.

As is well known, the conventional available magnesium hydroxides have a great strain, and their strain in the  $<101>$  direction is  $3.6 \times 10^{-3}$  at the least, and frequently reaches about  $10 \times 10^{-3}$ . The crystallite size of the conventional magnesium hydroxides is small, and about 700 Å at the largest, and generally in the range of 100 to 700 Å. Furthermore, they have a high surface area determined by the BET method, which is  $20 \text{ m}^2/\text{g}$  at the smallest, and in the range of 20 to 100  $\text{m}^2/\text{g}$ . The great strain in the structure of magnesium hydroxide means that the polarity of the surface of the crystallites is large, and the crystallites tend to aggregate secondarily with water as a binder. Thus, secondary aggregation occurs readily, and the crystallites aggregate into particles with a size of 10 to 100 microns. Thus, even after the drying of magnesium hydroxide, the aggregate contains negligible amounts of water and air trapped

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therin. Since the conventional available magnesium hydroxides have such a great strain in their structure as mentioned above, they have poor affinity for thermoplastic synthetic resins, especially those having great hydrophobicity or small polarity, for example, polyolefins. In addition, presumably because of the strong aggregation of its crystallites, its dispersibility in resins is extremely poor. In fact, resin compositions containing the conventional available magnesium hydroxides have poor shapability and exhibit reduced shaping efficiency. Furthermore, the poor affinity of the magnesium hydroxides with resins tends to result in the occurrence of spaces in the interface between the resin and the magnesium hydroxide particles. This, in turn, causes the deterioration of the physical properties of the resin composition, especially impact strength or elongation, and makes it difficult to disperse the magnesium hydroxide uniformly in the resin. The presence of water molecules and air formed as a result of the secondary aggregation of the crystallites, causes a poor appearance to the resulting fabricated articles, for example, by the formation of a silver pattern, presumably because they are released at the time of fabrication. Accordingly, the melt shaping of the resins is difficult to perform smoothly, and the dispersion of the magnesium hydroxide in the resin becomes nonuniform.

We have made extensive investigations in an attempt to overcome the difficulties associated with the structural characteristics of the conventional available magnesium hydroxides, and consequently found that a magnesium hydroxide having new structural characteristics clearly distinguishable from those of the conventional magnesium hydroxide can be provided, and that this magnesium hydroxide having the new structure has superior properties and is free from the disadvantages of the conventional magnesium hydroxides. It has also been found that the magnesium hydroxide having the new structure can be produced advantageously on a commercial scale by a very simple means which has not been performed heretofore.

Accordingly, it is an object of this invention to provide a magnesium hydroxide having a new structure and improved properties.

Another object of this invention is to provide a process for producing a magnesium hydroxide having a new structure with commercial advantage.

Still another object of this invention is to provide intermediates useful for providing a magnesium hydroxide having a new structure, and a process for preparing the intermediates.

Still another object of this invention is to provide a magnesium hydroxide coated with an anionic surface active agent which exhibits more favorably improved properties for use as a fire retardant for thermoplastic synthetic resins.

A further object of this invention is to provide a use of a magnesium hydroxide having a new structure which is coated or uncoated with an anionic surface active agent.

The above and other objects and advantages of this invention will become more apparent from the following description.

The magnesium hydroxide of this invention is expressed by  $Mg(OH)_2$ , and has a strain, in the,  $<101>$  direction, of not more than  $3.0 \times 10^{-3}$ . The magnesium hydroxide of this invention can be distinguished from the conventional magnesium hydroxides  $Mg(OH)_2$  in

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that the strain of the latter is at least  $3.6 \times 10^{-3}$ . Usually, the strain in the <101> direction of the magnesium hydroxide of this invention is within the range of, say,  $3.0 \times 10^{-3}$  to  $0.1 \times 10^{-3}$ . Furthermore, the magnesium hydroxide of this invention has a crystallite size in the <101> direction of more than 800 Å. This is another structural feature of the magnesium hydroxide of this invention since the crystallite sizes of the conventional magnesium hydroxides are 100 to 700 Å. Usually, the crystallite size of the magnesium hydroxide of this invention is within the range of more than 800 Å to 10000 Å.

Furthermore, while the conventional magnesium hydroxides have a specific surface area, determined by the BET method, of 20 to 100 m<sup>2</sup>/g, the specific surface area of the magnesium hydroxide of this invention determined by the same method is less than 20 m<sup>2</sup>/g, for example, at least 1 m<sup>2</sup>/g but less than 20 m<sup>2</sup>/g.

Magnesium hydroxides of this invention having optimum improved properties have a combination of the above three characteristic structural features.

The magnesium hydroxide of this invention having the new structure described above can be obtained by the hydrothermal treatment of basic magnesium chloride or magnesium nitrate of the following formula



wherein A is Cl or NO<sub>3</sub>, x is a number of more than 0 but less than 0.2, and m is a number of 0 to 6, which differs from the conventional magnesium hydroxides expressed by Mg(OH)<sub>2</sub> or the conventional magnesium hydroxychloride expressed by Mg(OH)Cl, in an aqueous medium at an elevated pressure.

The basic magnesium chloride or magnesium nitrate of the above formula can be formed by reacting magnesium chloride or magnesium nitrate with an alkaline substance in an aqueous medium, the amount of the alkaline substance being a controlled amount with respect to the magnesium chloride or magnesium nitrate, preferably 0.3 to 0.95 equivalent, more preferably 0.5 to 0.95 equivalent, per equivalent of the latter.

In the preparation of the compound of formula Mg(OH)<sub>2-x</sub>A<sub>x</sub>mH<sub>2</sub>O used in the manufacture of the magnesium hydroxide of this invention, it is preferred that the equivalent ratio of [OH] of the alkaline substance to [Mg<sup>2+</sup>] of the magnesium chloride or magnesium nitrate be maintained at  $2[\text{OH}]/[\text{Mg}^{2+}] = 0.3 - 0.95$ , and also a sufficient amount of a chlorine ion be present during the reaction. Favorable results can be obtained by adding the alkaline substance, for example, calcium hydroxide, in a controlled amount meeting the above equivalent relationship to an aqueous solution containing, for example, calcium chloride as well as magnesium chloride. Mg(OH)<sub>2</sub> obtained by conventional processes cannot become a magnesium hydroxide having the new structure described hereinabove even when it is heated in an aqueous medium at an elevated temperature. Basic magnesium nitrate can be prepared in the same way as above except that magnesium nitrate is used instead of magnesium chloride, and can equally be used in the preparation of the novel Mg(OH)<sub>2</sub> of this invention.

The reaction of forming basic magnesium chloride or magnesium nitrate is carried out at a temperature of about 0° to about 50° C, preferably at about 10° to about 20° C. The reaction is carried out in an aqueous medium under conditions such that the magnesium chloride or magnesium nitrate can be fully contacted with the alka-

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line substance. For example, this can be accomplished by adding calcium hydroxide in an amount to satisfy the above-mentioned equivalent relation to an aqueous solution of magnesium chloride, magnesium nitrate or both magnesium chloride and calcium chloride. Examples of the alkaline substance are calcium hydroxide, ammonia and alkali metal hydroxides.

The magnesium hydroxide having the novel structure in accordance with this invention can be produced by heating the basic magnesium chloride or basic magnesium nitrate Mg(OH)<sub>2-x</sub>A<sub>x</sub>mH<sub>2</sub>O formed by the method described above, in an aqueous medium at an elevated pressure, preferably at least about 5 Kg/cm<sup>2</sup>, for example, about 5 to 30 Kg/cm<sup>2</sup>. At this time, it is not necessary to isolate the basic magnesium chloride from the reaction mixture containing it, but the reaction mixture can be directly heated at an elevated pressure. This procedure is preferred. The magnesium hydroxide having the new structure in accordance with this invention cannot be formed even when ordinary magnesium hydroxide or known magnesium hydroxychloride is heated in the same way in an aqueous medium at an elevated pressure. The basic magnesium chloride or magnesium nitrate Mg(OH)<sub>2-x</sub>A<sub>x</sub>mH<sub>2</sub>O may be those which are prepared by other procedures. The heating treatment at an elevated pressure can be performed at a temperature of, say, about 150° to about 250° C.

As compared with the conventional magnesium hydroxides, the magnesium hydroxide having the new structure described hereinabove has a markedly small strain in the <101> direction, a large crystallite size in the same direction, and a very low specific surface area determined by the BET method. Because of these structural characteristics, the surface polarity of the crystallites is extremely small or nearly zero, and the secondary aggregation of the crystallites does not appreciably occur. In addition, the magnesium hydroxide of this invention is non-bulky, and has a low concentration of vacancy. This serves to overcome the disadvantages of the poor affinity of magnesium hydroxide with resins, poor shapability, or poor surface characteristics of the shaped articles which are associated with the incorporation of the conventional magnesium hydroxides in thermoplastic resins. Also, the defect of the deteriorated physical strength of the shaped articles can be eliminated.

The strain in the <101> direction, the crystallite size in the same direction, and the specific surface area by the BET method are measured by the following methods. Method of measuring the strain in the <101> direction and the crystallite size:

On the basis of the following expression,  $(\sin \theta/\lambda)$  is plotted in the axis of abscissas, and  $(\beta \cos \theta/\lambda)$  in the axis of ordinates. The crystallite size ( $\epsilon$ ) is sought from the inverse number of the intercept, and the strain ( $\eta$ ) is obtained by multiplying the gradient by  $\frac{1}{2}$ .

$$\frac{A \cos \theta}{\lambda} = \frac{1}{\epsilon} + 2\eta \frac{\sin \theta}{\lambda}$$

wherein  $\lambda$  is the wavelength of X-rays used (when they are Cu-K<sub>α</sub> rays, the wavelength is 1.542 Å);  $\theta$  is the Bragg's angle; and  $\beta$  is the true half-maximum breadth in radian.

In the above equation,  $\beta$  is determined by the following method.

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The diffraction profiles of the (101) and (202) planes are measured using Cu-K<sub>α</sub> rays generated at 35 KV and 15 mA as a source of X-rays. The measuring conditions are as follows:

Velocity of the goniometer:	1°/min.
Chart velocity:	10 mm/min.
Slit width:	(101) planes (202) planes
Divergence slit:	1°
Receiving slit:	0.3 mm
Scattering slit:	1° 2°

-continued

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ASTM No. 12-123- dA	1/I <sub>1</sub>	dA	1/I <sub>1</sub>
7.31	100	1.85	10
5.87	20	1.72	10
4.12	40	1.68	10
4.04	40	1.59	40
3.64	60	1.55	60
3.07	20	1.52	10
2.76	10	1.51	20
2.63	60	1.48	10
2.53	40	1.45	40
2.43	20		
2.40	100		
2.26	60		
2.24	40		

With regard to the profiles obtained, the width (B<sub>0</sub>) at a height half of the height from the background to the diffraction peak is measured. From the relation of the split width (8) of K<sub>α1</sub> and K<sub>α2</sub> to 2θ shown in FIG. 1 of the accompanying drawings, 8 of the (101) and (202) planes to 2θ is read. Then, on the basis of the B<sub>0</sub> and 8 values, B is determined from the relation between B/B<sub>0</sub> and B/B, shown in FIG. 2. The diffraction profiles of high purity (99.999%) silicon are measured at slit widths 1°-0.3 mm-1°, and the half-maximum breadth (b) is determined. This half-maximum breadth is plotted against 2θ to make a graphic representation showing the relation between b and 2θ. From b corresponding to 2θ in the (101) and (202) planes, b/B is determined, as shown in FIG. 3. β is determined from the relation between b/B and B/B. Method of measuring the specific surface area by the BET method:

Using a nitrogen adsorption isotherm method, the specific surface area is determined by a three-point plotting of nitrogen adsorption isotherm at -196° C. In the calculation, the cross sectional area of N<sub>2</sub> molecules adsorbed is set at 16.2 Å<sup>2</sup>. The specimens are subjected to the nitrogen adsorption test after having been evacuated in vacuum at 100° C for 30 minutes.

Known basic magnesium chlorides whose existence have been confirmed and which have been registered at ASTM are listed below. The numbers in the last column represent values for x.

ASTM No.	Formula	Mg(OH) <sub>2</sub> <sub>x</sub> Cl <sub>2</sub> <sub>m</sub> H <sub>2</sub> O	
12-116	Mg <sub>2</sub> Cl <sub>2</sub> (OH) <sub>4</sub> ·4H <sub>2</sub> O	2/3	
12-122	Mg <sub>2</sub> Cl <sub>2</sub> (OH) <sub>4</sub> ·4H <sub>2</sub> O	1/3	
12-123	Mg <sub>2</sub> Cl <sub>2</sub> (OH) <sub>4</sub> ·5H <sub>2</sub> O	1/5	
12-131	Mg <sub>2</sub> Cl <sub>2</sub> (OH) <sub>4</sub> ·4H <sub>2</sub> O	1/2	
7-403	Mg <sub>2</sub> Cl <sub>2</sub> (OH) <sub>4</sub> ·3H <sub>2</sub> O	1/2	
7-409	Mg <sub>2</sub> O·9Mg(OH) <sub>2</sub> ·5H <sub>2</sub> O	1/5	
7-412	Mg <sub>2</sub> Cl <sub>2</sub> (OH) <sub>4</sub> ·4H <sub>2</sub> O	1/2	
7-416	Mg <sub>2</sub> Cl <sub>2</sub> (OH) <sub>4</sub> ·3H <sub>2</sub> O	1/3	
7-420	Mg <sub>2</sub> Cl <sub>2</sub> (OH) <sub>4</sub> ·4H <sub>2</sub> O	1/3	
7-419	Mg <sub>2</sub> Cl <sub>2</sub> (OH) <sub>4</sub> ·Cl·2H <sub>2</sub> O	1/2	
3-0100	Mg <sub>2</sub> Cl <sub>2</sub>	1	
12-410	β-Mg <sub>2</sub> (OH) <sub>4</sub> Cl	1/2	
11-328	Mg <sub>2</sub> Cl <sub>2</sub>	1	
12-133	Mg <sub>2</sub> O <sub>2</sub> (OH) <sub>4</sub> ·2H <sub>2</sub> O	2/3	

As will be clear from the above list, ASTM No. 12-123 and No. 7-409 in which x is (1/5=0.2) smaller among those listed above are most similar to the basic magnesium chloride of this invention. The X-ray diffraction data of these known compounds described by ASTM and the X-ray diffraction data of the novel basic magnesium chloride of this invention measured by the same method are tabulated below.

ASTM No. 12-123- dA	1/I <sub>1</sub>	dA	1/I <sub>1</sub>
11.2	60	2.04	40
7.99	100	1.97	10

ASTM No. 7-409- dA	1/I <sub>1</sub>	hkl
11.1	25	002
7.93	100	100
7.27	55	102
5.87	20	103
4.11	30	104
4.03	25	202
3.68	6	106, 006
3.63	20	204
3.48	4	202
3.08	4	106
3.07	4	206
2.768	2	113
2.758	4	008
2.636	25	113
2.629	4	304
2.531	4	115
2.484	12	211
2.436	10	213
2.404	90	211
2.344	4	105
2.261	60	1010
2.144	14	
1.977	2	
1.888	2	
1.866	4	
1.852	2	
1.840	2	
1.803	2	
1.724	4	
1.709	3	
1.696	10	
1.626	1	
1.590	13	
1.575	2	
1.566	35	(20)
1.537	12	
1.529	8	
1.523	16	
1.507	2	

## Basic magnesium chloride of the invention:

Table 1

d(A)	1/I <sub>1</sub>	hkl
6.18	100	003
4.09	48	006
2.704	27	101
2.263	34	106
2.030	9	108
1.563	39	110
1.536	14	113

(Note) The hexagonal lattice constants are a<sub>0</sub> = 3.13 Å, c<sub>0</sub> = 24.6 Å.

It is clear from the above X-ray diffraction data that the magnesium chloride Mg(OH)<sub>2</sub><sub>x</sub>Cl<sub>2</sub><sub>m</sub>H<sub>2</sub>O of this invention has the novel structure different from the structures of the known compounds.

Similar data to those in Table 1 of the novel basic magnesium nitrate, an intermediate, of this invention represented by the formula Mg(OH)<sub>2</sub><sub>x</sub>(NO<sub>3</sub>)<sub>m</sub>H<sub>2</sub>O are shown in Table 2 below. The existence of basic magnesium nitrate is not described by ASTM.

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Table 2

d(Å)	L/I <sub>1</sub>	hkl
8.18	100	003
4.07	42	006
2.630	15	102
2.366	15	103
2.021	9	108
1.563	15	110
1.536	9	113

(Note) The hexagonal lattice constants are a<sub>0</sub> = 3.12 Å, c<sub>0</sub> = 24.4 Å.

If desired, the magnesium hydroxide of the invention may be treated with an anionic surface active agent to form solid particles of magnesium hydroxide coated with the surfactant. This form is more preferred in using it as a fire retardant or a fire-retarding filler for thermoplastic resins or water-soluble paints. The coating can be performed by contacting the magnesium hydroxide with anionic surfactants. For example, an aqueous solution of a desired amount of an anionic surfactant is mixed with solid particles of the magnesium hydroxide under conditions such that they contact each other sufficiently intimately, for example, by agitating them sufficiently, or by hydrothermal treatment at 120° to 250° C thereby to form a solid powder of magnesium hydroxide coated with the anionic surfactant. By this contacting operation, the surfactant is chemically adsorbed onto the surface of the solid particles of the magnesium hydroxide, and this can lead to more improved properties when the magnesium hydroxide is incorporated in thermoplastic synthetic resins or water-soluble paints.

The amount of the anionic surfactant to be coated can be adjusted optionally. Solid powder of the magnesium hydroxide of this invention coated by using an aqueous solution containing about 5 millimoles to about 30 millimoles, per liter of water, of the surfactant is preferred, for example. The amount of the anionic surfactant adsorbed onto the solid particles of the magnesium hydroxide of this invention is preferably about 1 to about 3 times, more preferably about 1 to about 2.5 times, the amount (X in millimoles) required to coat the entire surface of the solid particles (one gram) with a monolayer of the surfactant molecules. The amount X (millimoles) can be calculated in accordance with the following equation.

$$X = \frac{Y}{6.02 \times C} \quad (\text{millimoles})$$

wherein C is the absolute value of the adsorption cross-sectional area (Å<sup>2</sup>) per molecule of the anionic surfactant used, and Y is the absolute value of the specific surface area (m<sup>2</sup>/g) of the magnesium hydroxide of this invention.

According to this invention, there can be provided a composition containing uncoated magnesium hydroxide of this invention or the magnesium hydroxide of this invention coated with an anionic surface active agent. For example, compositions having improved properties, especially those useful for melt shaping, can be provided by incorporating the coated or uncoated magnesium hydroxide of this invention in a thermoplastic synthetic resin, preferably those having great hydrophobicity and great non-polarity, as a fire retardant or fire-retarding filler in an amount of about 50 to about 250 parts by weight per 100 parts by weight of the resin. Examples of the thermoplastic synthetic resin include styrene resins such as a homo- or copolymer of styrene, olefin resins such as homo- or copolymers of olefins,

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polyester resins, polycarbonate resins, nylon resins, acetal resins, and blends of these resins. These compositions may be provided in the form of melt-shaped articles. Furthermore, by incorporating the coated or uncoated magnesium hydroxide of this invention in paints or lacquers in an amount of about 5 to about 150 parts by weight per 100 parts by weight of the resin vehicle, paint compositions having improved properties can be obtained.

Various conventional additives may further be incorporated in the thermoplastic synthetic resin compositions or paint compositions in accordance with this invention.

Examples of these additives are coloring agents (organic and inorganic pigments) such as isoindolinone, cobalt aluminate, carbon black, or cadmium sulfide; other fillers such as calcium carbonate, alumina, zinc oxide or talc; antioxidants such as 2,6-di-t-butyl-4-methylphenol, 2,2'-methylenebis(4-methyl-6-t-butyl-phenol), diaryl thiadipropionate or tridecyl phosphite; ultraviolet absorbers such as 2-hydroxy-4-methoxy benzophenone, 2(2'-hydroxy-5'-methylphenyl) benzotriazole, 2-ethylhexyl-2-cyano-3,3-diphenyl acrylate, phenyl salicylate or nickel-bisocetyl phenyl sulfide; plasticizers such as di-2-ethyl hexyl phthalate, di-n-butyl phthalate, butyl stearate, or epoxidized soybean oil; and lubricants such as zinc stearate, calcium, aluminum and other metal soaps, or polyethylene wax.

These additives can be used in customary amounts. For example, the amount of the coloring agent is about 0.1 to about 3 parts by weight; the amount of the other filler is up to about 20 parts by weight; the amount of the antioxidant or ultraviolet absorber is about 0.001 to about 5 parts by weight; the amount of the plasticizer is up to about 20 parts by weight; and the amount of the lubricant is up to about 10 parts by weight. All these amounts are based on 100 parts by weight of the resin component.

The anionic surface active agent used to coat the magnesium hydroxide of this invention includes, for example, alkali metal salts of higher fatty acids of the formula



wherein R is an alkyl group containing 8 to 30 carbon atoms, and M is an alkali metal atom, alkyl sulfate salts of the formula



wherein R and M are the same as defined above, alkylsulfonate salts of the formula



wherein R and M are the same as defined above, alkylarylsulfonate salts of the formula



wherein R and M are the same as defined above, and sulfosuccinate ester salts of the formula



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wherein R and M are the same as defined above. These anionic surfactants can be used either alone or in admixture of two or more.

Specific examples of the surface active agent are sodium stearate, potassium behenate, sodium montanate, potassium stearate, sodium oleate, potassium oleate, sodium palmitate, potassium palmitate, sodium laurate, potassium laurate, sodium dilaurylbenzenesulfonate, potassium octadecylsulfate, sodium laurylsulfonate or disodium 2-sulfoethyl  $\alpha$ -sulfostearate.

The following Examples and Comparative Examples illustrate the present invention more specifically.

#### EXAMPLE 1

15 Five liters of an aqueous solution of magnesium chloride with a concentration of 1.5 moles/liter (the temperature of the solution was 15° C) was placed in a reaction vessel having a capacity of about 10 liters, and the solution was sufficiently stirred by means of a stirrer. An ammonia solution was with a concentration of 10 moles/liter (the temperature of the solution 15° C) was added in an amount of 1.35 liters (corresponding to 0.9 equivalent of the magnesium chloride, i.e. per equivalent of magnesium chloride) over the course of about 10 minutes.

20 A part of the resulting suspension was immediately filtered at reduced pressure, and then washed thoroughly with water and then with acetone. The product was dried for about 2 hours at room temperature, and analyzed by X-ray diffraction and by a chemical analysis method. By the X-ray diffraction, the product was identified as basic magnesium chloride having the structure defined in the present invention. The chemical analysis showed that this product had the composition  $Mg(OH)_{1.93}Cl_{0.07} \cdot mH_2O$ . The presence of water of crystallization was confirmed by DTA and TGA. Immediately after the reaction, a greater portion of the remaining suspension was placed in a 20-liter autoclave, and hydrothermally treated at 180° C for 8 hours. This heat-treatment was carried out within about 2 hours from the end of the reaction because this unstable substance had to be treated while it remained undecomposed. After the hydrothermal treatment, the product was filtered at reduced pressure, washed with water and dried. The product obtained was identified as magnesium hydroxide by X-ray diffraction. It had a strain in the <101> direction of  $0.970 \times 10^{-3}$ , a crystallite size in the <101> direction of 4200 Å, and a specific surface area by the BET method of  $6.7 \text{ m}^2/\text{g}$ .

#### EXAMPLE 2

25 The same procedure as in Example 1 was performed except that the ammonia solution was added in an amount of 1.05 liters (corresponding to 0.7 equivalent of magnesium chloride) over the course of about 7 minutes. A part of the resulting suspension was immediately filtered at reduced pressure, and then thoroughly washed with water and then with acetone. The resulting product was subjected to an X-ray diffraction analysis and a chemical analysis. By the X-ray diffraction, it was identified as the novel substance shown in Table 1. The chemical analysis showed that this product had the composition  $Mg(OH)_{1.92}Cl_{0.08} \cdot mH_2O$ .

30 On the other hand, the reaction mixture was placed in a 10-liter autoclave immediately after the reaction, and hydrothermally treated at 170° C for 8 hours. The product was filtered at reduced pressure, washed with water, and dried. The resulting magnesium hydroxide had

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a strain in the <101> direction of  $1.20 \times 10^{-3}$ , a crystallite size in the <101> direction of 5260 Å, and a specific surface area, by the BET method, of  $4.2 \text{ m}^2/\text{g}$ .

#### EXAMPLE 3

35 The same procedure as in Example 1 was performed except that 1.425 liters (corresponding to 0.95 equivalent of the magnesium chloride) of the ammonia solution was added over the course of about 10 minutes. A part of the reaction mixture was removed, and the remainder was immediately transferred to a 10-liter autoclave where it was hydrothermally treated at 200° C for 4 hours.

40 The reaction mixture previously removed, immediately after the reaction, was filtered at reduced pressure, and washed with water and then with acetone. The product was subjected to an X-ray diffraction analysis and a chemical analysis. The X-ray diffraction showed that the product was the novel substance shown in Table 1. The chemical analysis showed that the product had the composition  $Mg(OH)_{1.93}Cl_{0.07} \cdot mH_2O$ .

45 The hydrothermally treated product was filtered at reduced pressure, washed with water, and then dried.

50 The resulting magnesium hydroxide had a strain in the <101> direction of  $2.05 \times 10^{-3}$ , a crystallite size in the <101> direction of 2840 Å, and a specific surface area, by the BET method, of  $8.9 \text{ m}^2/\text{g}$ .

#### EXAMPLE 4

55 Ten liters of a mixed aqueous solution of magnesium chloride and calcium chloride (the by-product from the process of producing sodium chloride from sea water by an ion-exchange membrane method;  $Mg^{2+} = 1.98 \text{ mole/liter}$ ,  $Ca^{2+} = 0.765 \text{ mole/liter}$ ) and 8.2 liters (corresponding to 0.8 equivalent of magnesium chloride) of an aqueous solution of calcium hydroxide in a concentration of 1.54 mole/liter were each maintained at 30° C. One liter of water was placed in a 2-liter reactor equipped with an overflowing device, and stirred by a stirrer. The temperature of the water was adjusted to 30° C. Using metering pumps, the mixed aqueous solution of magnesium chloride and calcium chloride and the aqueous solution of calcium hydroxide were fed into the reactor at a feed rate of 100 mL/min., and 82 mL/min., respectively to perform the reaction. After the reaction, 16 liters of the resulting suspension was immediately transferred to a 30-liter autoclave, and hydrothermally treated at 145° C for 8 hours. The remainder of the reaction mixture was filtered at reduced pressure, washed with water and acetone, and dried at room temperature for 4 hours. The product was subjected to an X-ray diffraction analysis and a chemical analysis. The X-ray diffraction showed that the product was the novel substance shown in Table 1. As a result of the chemical analysis, this product was found to have the composition  $Mg(OH)_{1.93}Cl_{0.07} \cdot mH_2O$ . The hydrothermally treated product was filtered at reduced pressure, washed with water, and dried. The magnesium hydroxide so obtained had a strain in the <101> direction of  $1.80 \times 10^{-3}$ , a crystallite size in the <101> direction of 2250 Å, and a specific surface area by the BET method of  $12.7 \text{ m}^2/\text{g}$ .

#### EXAMPLE 5

60 Two liters of an aqueous solution of magnesium nitrate in a concentration of 2 mole/liter (the temperature of the solution was 15° C) was placed in a reactor with a capacity of about 5 liters, and stirred thoroughly by a

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wherein R and M are the same as defined above. These anionic surfactants can be used either alone or in admixture of two or more.

Specific examples of the surface active agent are sodium stearate, potassium behenate, sodium montanate, potassium stearate, sodium oleate, potassium oleate, sodium palmitate, potassium palmitate, sodium laurate, potassium laurate, sodium dilaurylbenzenesulfonate, potassium octadecylsulfate, sodium laurylsulfonate or disodium 2-sulfoethyl  $\alpha$ -sulfostearate.

The following Examples and Comparative Examples illustrate the present invention more specifically.

#### EXAMPLE 1

15 Five liters of an aqueous solution of magnesium chloride with a concentration of 1.5 moles/liter (the temperature of the solution was 15° C) was placed in a reaction vessel having a capacity of about 10 liters, and the solution was sufficiently stirred by means of a stirrer. An ammonia solution was with a concentration of 10 moles/liter (the temperature of the solution 15° C) was added in an amount of 1.35 liters (corresponding to 0.9 equivalent of the magnesium chloride, i.e. per equivalent of magnesium chloride) over the course of about 10 minutes.

A part of the resulting suspension was immediately filtered at reduced pressure, and then washed thoroughly with water and then with acetone. The product was dried for about 2 hours at room temperature, and analyzed by X-ray diffraction and by a chemical analysis method. By the X-ray diffraction, the product was identified as basic magnesium chloride having the structure defined in the present invention. The chemical analysis showed that this product had the composition  $Mg(OH)_{1.59}Cl_{0.07}.mH_2O$ . The presence of water of crystallization was confirmed by DTA and TGA. Immediately after the reaction, a greater portion of the remaining suspension was placed in a 20-liter autoclave, and hydrothermally treated at 180° C for 8 hours. This heat-treatment was carried out within about 2 hours from the end of the reaction because this unstable substance had to be treated while it remained undecomposed. After the hydrothermal treatment, the product was filtered at reduced pressure, washed with water and dried. The product obtained was identified as magnesium hydroxide by X-ray diffraction. It had a strain in the <101> direction of  $0.970 \times 10^{-3}$ , a crystallite size in the <101> direction of 4200 Å, and a specific surface area by the BET method of  $6.7 \text{ m}^2/\text{g}$ .

#### EXAMPLE 2

The same procedure as in Example 1 was performed except that the ammonia solution was added in an amount of 1.05 liters (corresponding to 0.7 equivalent of magnesium chloride) over the course of about 7 minutes. A part of the resulting suspension was immediately filtered at reduced pressure, and then thoroughly washed with water and then with acetone. The resulting product was subjected to an X-ray diffraction analysis and a chemical analysis. By the X-ray diffraction, it was identified as the novel substance shown in Table 1. The chemical analysis showed that this product had the composition  $Mg(OH)_{1.49}Cl_{0.10}.mH_2O$ .

On the other hand, the reaction mixture was placed in a 10-liter autoclave immediately after the reaction, and hydrothermally treated at 170° C for 8 hours. The product was filtered at reduced pressure, washed with water, and dried. The resulting magnesium hydroxide had

5 a strain in the <101> direction of  $1.20 \times 10^{-3}$ , a crystallite size in the <101> direction of 5260 Å, and a specific surface area, by the BET method, of  $4.2 \text{ m}^2/\text{g}$ .

#### EXAMPLE 3

10 The same procedure as in Example 1 was performed except that 1.425 liters (corresponding to 0.95 equivalent of the magnesium chloride) of the ammonia solution was added over the course of about 10 minutes. A part of the reaction mixture was removed, and the remainder was immediately transferred to a 10-liter autoclave where it was hydrothermally treated at 200° C for 4 hours.

15 The reaction mixture previously removed, immediately after the reaction, was filtered at reduced pressure, and washed with water and then with acetone. The product was subjected to an X-ray diffraction analysis and a chemical analysis. The X-ray diffraction showed that the product was the novel substance shown in Table 1. The chemical analysis showed that the product had the composition  $Mg(OH)_{1.59}Cl_{0.07}.mH_2O$ .

20 The hydrothermally treated product was filtered at reduced pressure, washed with water, and then dried.

25 The resulting magnesium hydroxide had a strain in the <101> direction of  $2.05 \times 10^{-3}$ , a crystallite size in the <101> direction of 2840 Å, and a specific surface area, by the BET method, of  $8.9 \text{ m}^2/\text{g}$ .

#### EXAMPLE 4

30 Ten liters of a mixed aqueous solution of magnesium chloride and calcium chloride (the by-product from the process of producing sodium chloride from sea water by an ion-exchange membrane method;  $Mg^{2+} = 1.58$  mole/liter,  $Ca^{2+} = 0.765$  mole/liter) and 8.2 liters (corresponding to 0.8 equivalent of magnesium chloride) of an aqueous solution of calcium hydroxide in a concentration of 1.54 mole/liter were each maintained at 30° C. One liter of water was placed in a 2-liter reactor equipped with an overflowing device, and stirred by a stirrer. The temperature of the water was adjusted to 30° C. Using metering pumps, the mixed aqueous solution of magnesium chloride and calcium chloride and the aqueous solution of calcium hydroxide were fed into the reactor at a feed rate of 100 ml/min., and 82 ml/min., respectively to perform the reaction. After the reaction, 16 liters of the resulting suspension was immediately transferred to a 30-liter autoclave, and hydrothermally treated at 145° C for 8 hours. The remainder of the reaction mixture was filtered at reduced pressure, 35 washed with water and acetone, and dried at room temperature for 4 hours. The product was subjected to an X-ray diffraction analysis and a chemical analysis. The X-ray diffraction showed that the product was the novel substance shown in Table 1. As a result of the chemical analysis, this product was found to have the composition  $Mg(OH)_{1.59}Cl_{0.07}.mH_2O$ . The hydrothermally treated product was filtered at reduced pressure, washed with water, and dried. The magnesium hydroxide so obtained had a strain in the <101> direction of  $1.80 \times 10^{-3}$ , a crystallite size in the <101> direction of 2250 Å, and a specific surface area by the BET method of  $12.7 \text{ m}^2/\text{g}$ .

#### EXAMPLE 5

40 Two liters of an aqueous solution of magnesium nitrate in a concentration of 2 mole/liter (the temperature of the solution was 15° C) was placed in a reactor with a capacity of about 5 liters, and stirred thoroughly by a

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stirrer. Ammonia solution (at 15° C) in a concentration of 4 mole/liter was added in an amount of 1.8 liters (corresponding to 0.9 equivalent of the magnesium nitrate) over the course of about 20 minutes. Two liters of the resulting suspension was immediately transferred to a 5-liter autoclave, and hydrothermally treated at 170° C for 4 hours. The remainder (1.8 liters), immediately after the reaction, was filtered at reduced pressure, and thoroughly washed with acetone. The product was subjected to an X-ray diffraction analysis and a chemical analysis. The product was identified by the X-ray diffraction as the novel substance shown in Table 2. The chemical analysis showed that the product had the composition  $Mg(OH)_{1.17}(NO_3)_{0.17} \cdot mH_2O$ . The hydrothermally treated product was filtered at reduced pressure, 15 washed with water, and dried. The magnesium hydroxide so obtained had a strain in the <101> direction of  $2.40 \times 10^{-3}$ , a crystallite size in the <101> direction of 4200 Å, and a specific surface area by the BET method of 9.6 m<sup>2</sup>/g.

#### COMPARATIVE EXAMPLE 1

Two liters of an aqueous solution of magnesium chloride in a concentration of 1.5 moles/liter was maintained at 40° C, and thoroughly stirred. An aqueous 25 solution of calcium hydroxide in a concentration of 1.5 moles/liter was added to the stirred magnesium chloride solution in an amount of 2 liters (corresponding to one equivalent of the magnesium chloride) over the course of about 60 minutes. The reaction mixture obtained was filtered at reduced pressure, and washed with water. The product was filtered, and dried at 80° C for 10 hours. The resulting product was identified as magnesium hydroxide by an X-ray diffraction analysis. 30 The product washed with water was suspended in 6 liters of water, and hydrothermally treated at 250° C for 8 hours in a 10-liter autoclave. The hydrothermally treated product was filtered at reduced pressure, washed with water, and dried. The product obtained had a strain in the <101> direction of  $3.70 \times 10^{-3}$ , a 35 crystallite size in the <101> direction of 568 Å, and a specific surface area by the BET method of 32 m<sup>2</sup>/g. The compound before the hydrothermal treatment had a strain in the <101> direction of  $4.76 \times 10^{-3}$ , a crystallite size in the <101> direction of 549 Å, and a 40 specific surface area by the BET method of 21 m<sup>2</sup>/g.

#### COMPARATIVE EXAMPLE 2

Four liters of an aqueous solution of magnesium chloride in a concentration of 1.5 moles/liter, and 4 liters of 50 an aqueous solution of calcium hydroxide in a concentration of 2.0 moles/liter were each maintained at 20° C. 500 ml of water was placed in a 1.5-liter reactor equipped with an overflowing device, and thoroughly stirred. Using metering pumps, the above aqueous solutions were each fed into the reactor at a rate of 40 ml/min. The amount of the alkali fed was adjusted to one equivalent per equivalent of magnesium chloride. In about 100 minutes, the reaction ended. A part of the resulting suspension was filtered at reduced pressure, 55 and washed with water and acetone. The product was analyzed by X-ray diffraction and found to be magnesium hydroxide. The remainder of the suspension, im-

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mediately after the reaction, was transferred to a 10-liter autoclave, and hydrothermally treated at 170° C for 8 hours. The hydrothermally treated product was filtered at reduced pressure, washed with water, and dried. The product had a strain in the <101> direction of  $3.70 \times 10^{-3}$ , a crystallite particle size in the <101> direction of 647 Å, and a specific surface area by the BET method of 26 m<sup>2</sup>/g. The compound before the hydrothermal treatment had a strain in the <101> direction of  $4.83 \times 10^{-3}$ , a crystallite particle size in the <101> direction of 476 Å, and a specific surface area by the BET method of 31 m<sup>2</sup>/g.

#### EXAMPLE 6

2.2 Kg of the magnesium hydroxide obtained in Example 2 having a strain of  $1.20 \times 10^{-3}$ , a crystallite particle size of 5260 Å, and a specific surface area of 4.2 m<sup>2</sup>/g was further dried at 150° C for 3 hours, and mixed with 1.8 Kg of polypropylene having a melt index of 6.0 and a density of 0.91 by a Henschel mixer. The composition was heated to about 230° C, and melt-kneaded in an extruder. The resulting resin composition was injection-molded into a plate. The physical properties and fire retardancy of the plate were evaluated by the ASTM standards and the UL standards. The results obtained are shown in Table 3.

#### EXAMPLE 7

2.2 Kg of the magnesium hydroxide obtained in Example 1 was placed in 10 liters of an aqueous solution of sodium stearate in a concentration of  $10 \times 10^{-3}$  mole/liter. The mixture was maintained at 80° C for 2 hours with stirring to coat the surface of the magnesium hydroxide with the sodium stearate. The product was filtered at reduced pressure, washed with water, and dried. Using the resulting dried product instead of the magnesium hydroxide used in Example 6, the same procedure as in Example 6 was performed. The results are shown in Table 3.

#### COMPARATIVE EXAMPLE 3

The same procedure as in Example 6 was performed except that 2.2 Kg of the magnesium hydroxide obtained in Comparative Example 1 having a strain of  $4.76 \times 10^{-3}$ , a crystallite size of 549 Å, and a specific surface area of 21 m<sup>2</sup>/g was used instead of the magnesium hydroxide used in Example 6. The results are shown in Table 3.

#### COMPARATIVE EXAMPLE 4

The same procedure as in Example 6 was performed except that the magnesium hydroxide obtained in Comparative Example 2 having a strain of  $3.70 \times 10^{-3}$ , a crystallite particle size of 647 Å, and a specific surface area of 26 m<sup>2</sup>/g was used instead of the magnesium hydroxide used in Example 6. The results are shown in Table 3.

#### CONTROL EXAMPLE 1

The polypropylene alone used in Example 6 was molded in the same way as in Example 6. The results are shown in Table 3.

Table 3

Runs	Injection pressure (#1)	Appearance of the surface of the shaped article	Iced impact strength (#2)	Tensile elongation (#3)	Fire retardancy (#4)
Example 6	105	No silver pattern	3.9	9.7	V-0

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Table 3-continued

Runs	Injection pressure (*1)	Appearance of the surface of the shaped article	Izod impact strength (*2)	Tensile elongation (*3)	Fire retardancy (*4)
Example 7	96	No silver pattern	5.2	21.0	V-O
Comparative 18D	18D	Silver pattern	0.8	1.9	HB
Example 3		present			
Comparative 14G	14G	Silver pattern	1.1	2.1	HB
Example 4		present			
Control	10D	No silver pattern	7.1	>100	Combustible
Example 1					

(\*1) Relative values of the injection pressure based on that used in molding polypropylene alone.

(\*2) ASTM D256 (Kg-cm/cm)

(\*3) ASTM D638 (S)

(\*4) UL Standard 94 V-E

## EXAMPLES 8 TO 15 AND CONTROL EXAMPLES 2 TO 5

The same procedure as in Example 6 was performed except that 100 parts by weight of each of the thermoplastic resins shown in Table 4 was blended with each of the uncoated magnesium hydroxide used in Example 6 and the magnesium hydroxide coated in the same manner as in Example 7 with each of the anionic surfactants shown in Table 4, and each of the resulting compositions was injection-molded at the pressures and temperatures shown in Table 4. The results obtained are shown in Table 4.

15 than  $3.0 \times 10^{-3}$ , (ii) a crystallite size in the  $<101>$  direction of more than 800 Å, and (iii) a specific surface area, determined by the BET method, of less than 20 m<sup>2</sup>/g.

20 3. The solid particles of magnesium hydroxide according to claim 1 wherein said anionic surface active agent is a member selected from the group consisting of alkali metal salts of higher fatty acids of the formula



25 wherein R represents an alkyl group containing 8 to 30 carbon atoms, and M represents an alkali metal

Table 4

Run	Resin	Amount of magnesium hydroxide (parts by weight per 100 parts by weight of resin)	Surface active agent		Molding temperature (°C)	Injection pressure (*1)	Appearance of the molded article	Izod impact strength (*2)	Tensile elongation (*3)	Fire retardancy (*4)
			Compound	Amount coated (% by weight based on Mg(OH) <sub>2</sub> )						
Ex. 8	Polyethylene	uncoated (135)			230-240	100	No silver pattern	2.1	18.1	V-O
Ex. 9		coated (135)	Sodium oleate	4.0	"	80	No silver pattern	3.4	29.5	V-O
Con. 2		Mg(OH) <sub>2</sub> not added			"	100	No silver pattern	1.0	>500	—
Ex. 10	Poly-styrene	uncoated (125)			240-250	100	No silver pattern	2.2	1.9	V-O
Ex. 11		coated (125)	Sodium lauryl benzene sulfonate	6.5	"	87	No silver pattern	4.1	2.6	V-O
Con. 3		Mg(OH) <sub>2</sub> not added			"	100	No silver pattern	7.0	35.0	—
Ex. 12	ABS (*5)	uncoated (125)			250-260	110	No silver pattern	18	2.9	V-O
Ex. 13		coated (125)	Sodium behenate	9.2	"	90	No silver pattern	21	4.1	V-O
Con. 4		Mg(OH) <sub>2</sub> not added			"	100	No silver pattern	40	50	—
Ex. 14	Nylon 6	uncoated (120)			250-300	110	Silver pattern present	8.3	4.2	V-O
Ex. 15		coated (120)	Sodium montanate	8.7	"	92	No silver pattern	14.1	6.0	V-O
Con. 5		Mg(OH) <sub>2</sub> not added			"	100	No silver pattern	19.0	65	—

Notes (\*1) to (\*4) are the same as the foregoing to Table 3.

(\*5) ABS is an acrylonitrile/butadiene/styrene copolymer with a weight ratio of 20/23/57.

What we claim is:

1. Solid particles of a magnesium hydroxide coated with an anionic surface active agent, said magnesium hydroxide having (i) a strain in the  $<101>$  direction of not more than  $3.0 \times 10^{-3}$ , (ii) a crystallite size in the  $<101>$  direction of more than 800 Å, and (iii) a specific surface area, determined by the BET method, of less than 20 m<sup>2</sup>/g.

2. A thermoplastic synthetic resin composition comprising 100 parts by weight of a thermoplastic synthetic resin and about 50 to about 250 parts by weight of a magnesium hydroxide coated or uncoated with an anionic surface active agent, said magnesium hydroxide having (i) a strain in the  $<101>$  direction of not more

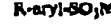
55 atom, alkylsulfate salts of the formula



wherein R and M are the same as defined above, alkylsulfonate salts of the formula



wherein R and M are the same as defined above, alkylarylsulfonate salts of the formula



4,098,762

15

16

wherein R and M are the same as defined above, and

sulfoguccinate ester salts of the formula



5 wherein R and M are the same as defined above.

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## Fusabond®

polymer modifiers

### **Fusabond® Resins as Coupling Agents in W & C Applications**

*Fusabond®* maleic anhydride grafted resins can be used as coupling agents between polymers -- mainly polyolefins such as polyethylene and polypropylene -- and fillers, in order to increase the filler acceptability of polymers. Typical levels of *Fusabond®* are in the range of 2-5 weight % based on the entire compound. Fillers that show an affinity to *Fusabond®* maleic anhydride grafted resins include flame-retardant fillers, such as alumina trihydrate (ATH) and magnesium hydroxide (Mg[OH]2). Improvement in properties also can be seen when *Fusabond®* is used in combination with common fillers such as calcium carbonate.

Extensive trials have been undertaken using *Fusabond®* E as a coupling agent in halogen-free, flame retardant wire and cable formulations containing approximately 65% ATH in a LLDPE/EVA matrix.

Figure 1 compares typical property data for extruded sheet made from compounds containing varying levels of *Fusabond®*, versus a control compound containing a silane-based coupling agent. Result show that *Fusabond®* has a significant influence on elongation at break and, therefore, the flexibility of a compound. Compared with liquid silanes, *Fusabond®* resins are solid particles, which generally can be handled more easily and dosed more accurately.

Figure 2 shows how heat-aging of the described PE/ATH formulations affects tensile strength, elongation at break, and viscosity (melt flow index).

In a related application, compounds containing polypropylene with approximately 65% calcium carbonate were examined. Figure 3 is a photomicrograph comparing the microstructures of compounds made with and without *Fusabond®* coupling agent. With coupling agent added at 4% weight, based on the entire compound weight, properties mainly impacted are stiffness, tensile strength and impact strength as shown in Figure 4. Note that impact strength is significantly improved (by almost a factor of 3) at temperatures of -20°C (-4°F) and 4°C (39°F), in the compounds containing *Fusabond®*.

**Figure 1.**

DuPont Industrial Polymers: Fusabond®

Page 2 of 3

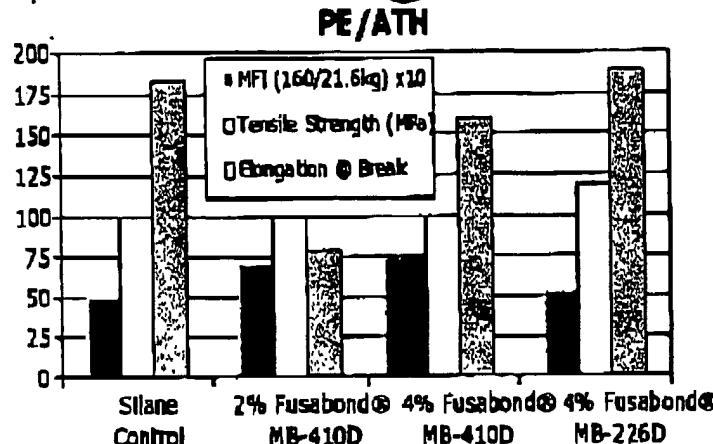


Figure 2.

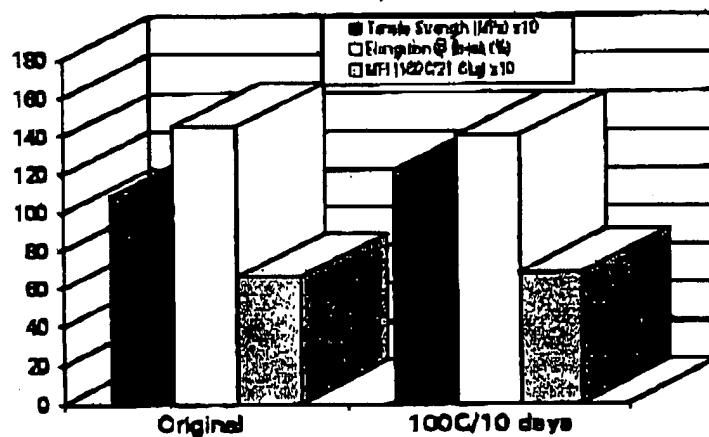
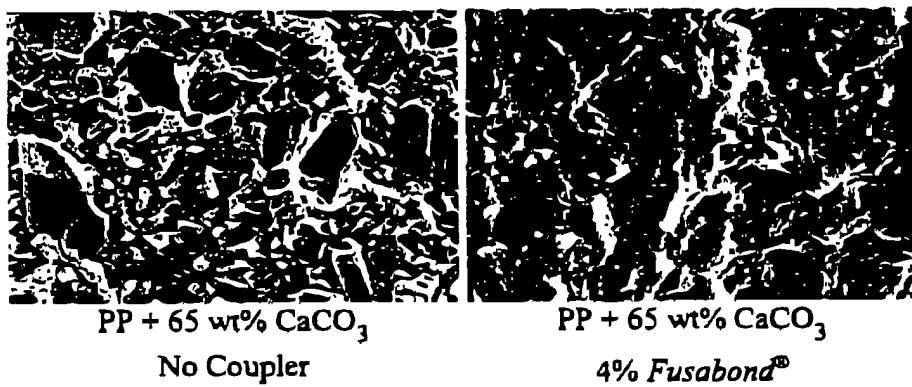
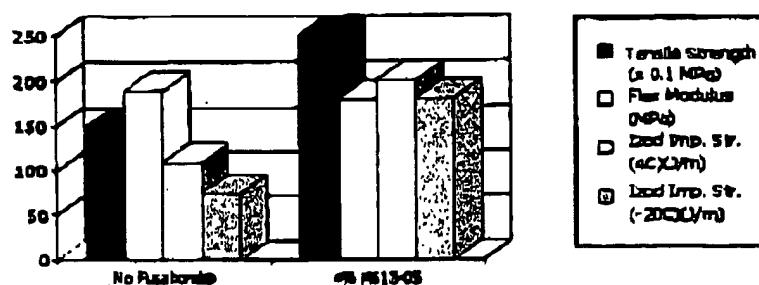
**PE/ATH Aging Performance****Figure 3. Effect of PP-g MAH (Fusabond® M613-05 on Fracture Toughness of PP/CaCO<sub>3</sub>)**

Figure 4.



The above examples show that *Fusabond®* can be used as a coupling agent for flame retardant fillers in custom wire and cable formulations.



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The technical data contained herein are guides to the use of DuPont resins. The advice contained herein is based upon tests and information believed to be reliable, but users should not rely upon it absolutely for specific applications because performance properties will vary with processing conditions. It is given and accepted at user's risk and confirmation of its validity and suitability in particular cases should be obtained independently. The DuPont Company makes no guarantees of results and assumes no obligations or liability in connection with its advice. This publication is not to be taken as a license to operate under, or recommendation to infringe, any patents.

**CAUTION:** Do not use in medical applications involving permanent implantation in the human body. For other medical applications, see "DuPont Medical Caution Statement", H-50102.

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**FACSIMILE TRANSMITTAL****TO**

**Name:** Examiner J. Gray  
**Firm:** United States Patent & Trademark  
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**Phone No.:** (703) 308-2381  
**Date:** May 13, 2003  
**Subject:** Application No. 09/815,311  
In re Application of: Eduardo GRIZANTE REDONDO et al.  
Filed March 23, 2001

**FROM**

**Name:** Gordon Klancnik  
**Phone No.:** (202) 408-4322  
**Fax # Verified by:** GPK (MD 905)  
**# Pages (incl. this):** 23  
**Our File No.:** 05788.0157

**Confirmation Copy to Follow: No****Message:**

Examiner Gray:

Thank you for your time on May 12, 2003. Further to our telephone conversations, Applicants re-submit herewith a copy of the date-stamped postcard and the Response After Final Under 37 C.F.R. § 1.116 dated January 24, 2003. It is the understanding of Applicants' representative that the Office has misplaced the previously filed Response. Applicants respectfully request that the Examiner correct the record to reflect that the Amendment had been filed with the Office on January 24, 2003.

Applicants presently expect to file a Notice of Appeal on May 14, 2003, to maintain pendency of the application while the Examiner has the opportunity to consider the filing. Accordingly, Applicants respectfully request an expeditious examination and reconsideration in light of these facts.

Respectfully submitted,



Gordon Klancnik  
Reg. No. 50,964

Dated: May 13, 2003

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AJS/AAH/GPK

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In Re Application of: Eduardo GRIZANTE REDONDO et al.

Application No.: 09/815,311

Group Art Unit: 1774

Filed: March 23, 2001

Examiner: J. Gray

For: LOW-SMOKE SELF-EXTINGUISHING ELECTRICAL CABLE AND FLAME-  
RETARDANT COMPOSITION USED THEREIN

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1. Amendment to Office Action dated November 14, 2002 (7 pages)
2. Attachments (U.S. Patent No. 4,098,762; and Dupont Industrial Polymers: *Fusabond*®)

Dated: January 24, 2003

Docket No.: 05788.0157-00

AJS/AAH/Gordon P. Klancnik - M.Beach at MD905



(Due Date: February 14, 2003)

*05788.0157-00*